ATROPINE SULFATE- atropine sulfate injection, solution
MWI

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Atropine Sulfate Injection

NDC 13985-354-04
VETone

ATROPINE SULFATE
Injection 1/120 Grain

Sterile Multiple Dose Vial

Keep out of reach of children.

For veterinary use only.

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

V1 510221 Net Contents: 100 mL

For Intravenous, Intramuscular, or Subcutaneous Use
COMPOSITION: **Each mL contains**:
Atropine Sulfate ..... 0.54 mg
Sodium Chloride ..... 9 mg
Benzyl Alcohol (preservative)..... 1.5%
Water for Injection ..... q.s.
pH adjusted with sulfuric acid when necessary.

DOSAGE AND ADMINISTRATION:
Dogs and Cats: Inject 1 mL for each 20 lbs. of body weight as a pre-anesthetic adjuvant, or to reduce salivation, bronchial secretions, or internal peristalsis associated with colic or diarrhea.

As an antidote for parasympathomimetic drugs, 1 mL for each 7.5 lbs. of body weight. It is suggested that 1/4 of the dosage be injected intravenous and the remainder intramuscular or subcutaneous.

WARNING: Poisonous alkaloid. Keep out of reach of children
Antidotes: warmth, emetics, cholinergics.

STORAGE: Store at room temperature between **15° - 30°C (59°- 86°F)**

Rev. 02/14

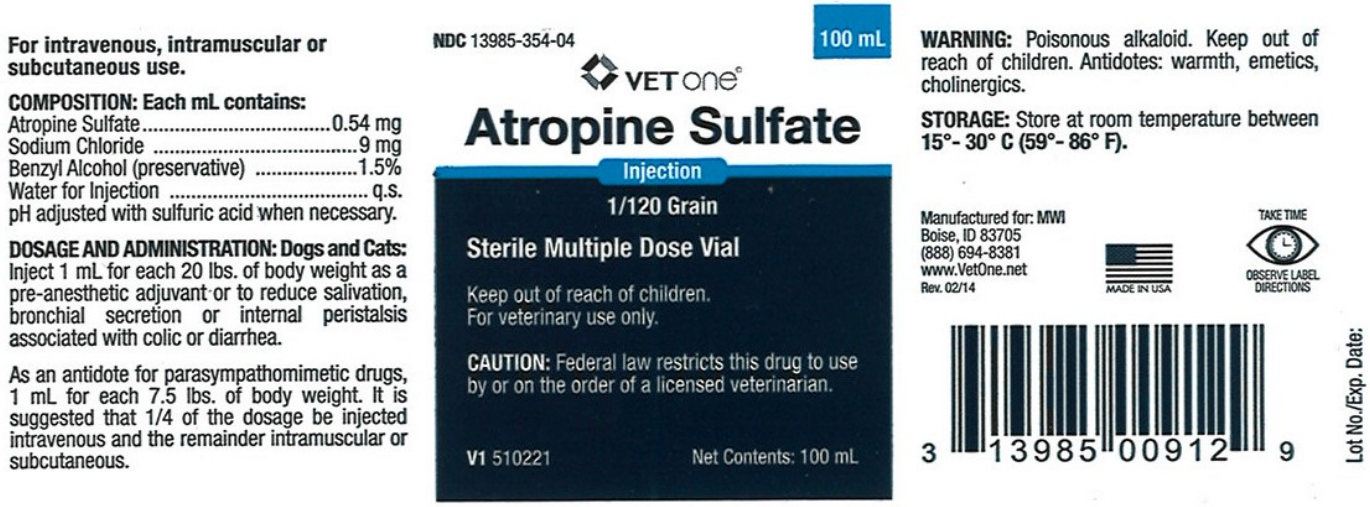
Manufactured for: **MWI**
Boise, ID 83705

(888) 694-8381

www.VetOne.net

TAKE TIME OBSERVE LABEL DIRECTIONS

Lot No./Exp. Date:



ATROPINE SULFATE

atropine sulfate injection, solution

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:13985-354
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ATROPINE SULFATE (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	0.54 mg in 1 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13985-354-04	100 mL in 1 VIAL, MULTI-DOSE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2008	

Labeler - MWI (019926120)

Revised: 6/2018 MWI